Supplemental Material

Disruption of Thyroid Hormone Receptor-Mediated Transcription and

Thyroid Hormone-Induced Purkinje Cell Dendrite Arborization by

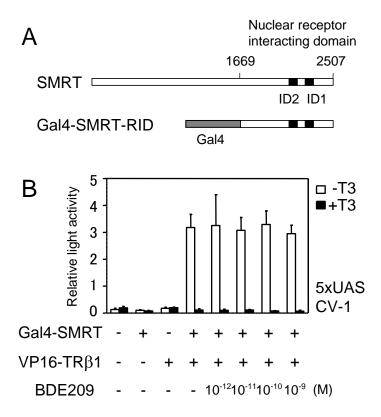
Polybrominated Diphenylethers.

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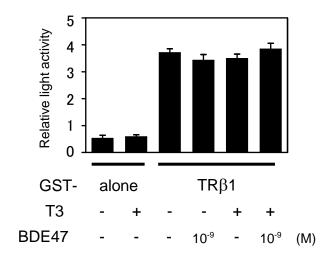
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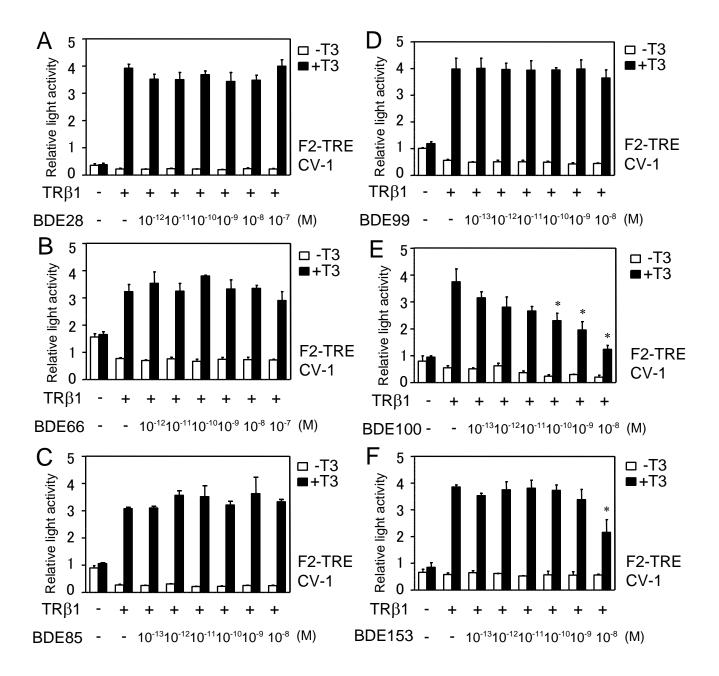


Effect of PBDE on TR-SMRT binding in the presence of T₃ in CV-1 cells.

(A) Schematic diagram of Gal4-SMRT. Gal4-SMRT contains amino acid residues 1669-2507 SMRT. (B) PBDE did not recruit SMRT to liganded TR. Expression plasmids harboring Gal4-DBD-fused SMRT (100 ng) were cotransfected with VP16-TR β 1-LBD (50 ng) and 5x UAS-TK-LUC (100 ng) into CV-1 cells. Cells were incubated with or without T₃ (10⁻⁷ M) and indicated concentrations of BDE209. Data represent mean \pm S.E.M. of experiments performed in triplicate. No statistical significance was determined.

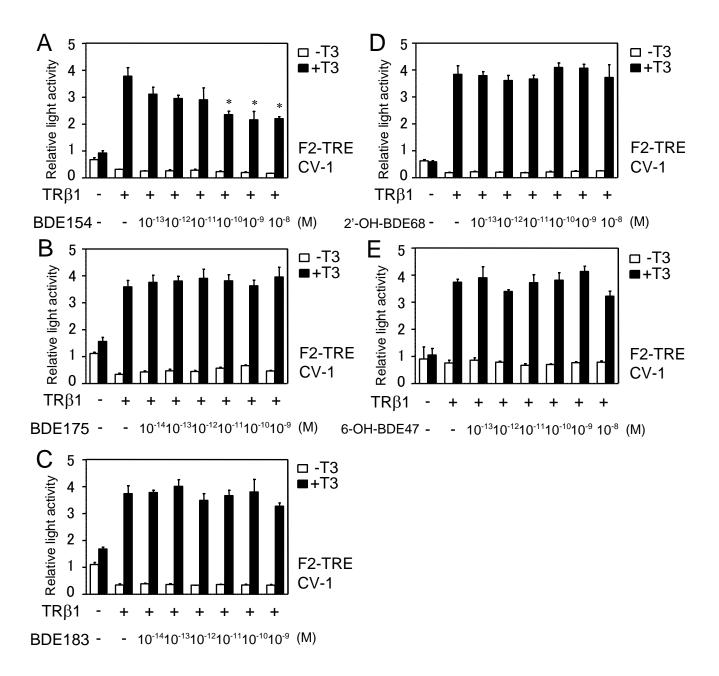


BDE47 did not alter the TR-TRE binding. GST-TR β 1 bound to sepharose beads is incubated with DIG-labeled-F2-TRE in protein-DNA binding buffer with or without T₃ (10⁻⁶ M) and BDE47 (10⁻⁹ M). Data represent mean \pm S.E.M. of experiments performed in triplicate. No statistical significance was uncovered.



Congener-specific suppression of TR-mediated transcription by PBDE.

(A-F) Expression plasmids encoding TR β 1 (10 ng) were cotransfected with F2-TK-LUC (100 ng) into CV-1 cells. Cells were incubated with or without 10⁻⁷ M T₃ and indicated amounts of listed PBDE. Data represent mean \pm S.E.M. of experiments performed in triplicate. p < 0.01 by ANOVA, compared with TR β 1 (+), T₃ (+), and PBDE (-).



Congener-specific suppression of TR-mediated transcription by PBDE/OH-PBDE.

(A-E) Expression plasmids encoding TR β 1 (10 ng) were cotransfected with F2-TK-LUC (100 ng) into CV-1 cells. Cells were incubated with or without 10^{-7} M T₃ and indicated amounts of listed PBDE/OH-PBDE. Data represent mean \pm S.E.M. of experiments performed in triplicate. p < 0.01 by ANOVA, compared with TR β 1 (+), T₃ (+), and PBDE/OH-PBDE (-).